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Substitute for form 1449A/B/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)				Complete if Known	
				Application Number	Unassigned
				Filing Date	March 3, 2004
				First Named Inventor	GRIFFITH et al.
				Group Art Unit	Unassigned
				Examiner Name	Unassigned
Sheet	1	of	1	Attorney Docket Number	227540

U.S. PATENT DOCUMENTS						
Examiner Initials	Doc. No.	U.S. Patent Document		Name of Patentee or Applicant	Date of Publication	Filing Date If Appropriate
		Application or Patent Number	Kind Code			
APW	AA	6,040,138		Lockhart et al.	Mar. 21, 2000	Sep. 15, 1995
APW	AB	6,197,506	B1	Fodor et al.	Mar. 6, 2001	Apr. 8, 1998
APW	AC	6,485,908	B1	Petit et al.	Nov. 26, 2002	Aug. 14, 1998

FOREIGN PATENT DOCUMENTS								
Examiner Initials	Doc. No.	Foreign Patent Document			Name of Patentee or Applicant	Date of Publication	Translation	
		Office	Application or Patent Number	Kind Code			Yes	No**
APW	AD	WO	98/00014	A1	Regents of the University of California	Jan. 8, 1998		
APW	AE	WO	99/09210	A2	Institut Pasteur	Feb. 25, 1999		

OTHER - NON PATENT LITERATURE DOCUMENTS				
Examiner Initials	Doc. No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number (s), publisher, city and/or country where published.	Translation	
			Yes	No**
APW	AF	COREY et al., "Ionic Basis of the Receptor Potential in a Vertebrate Hair Cell," <i>Nature</i> , Vol. 281, pp. 675-677 (1979)		
APW	AG	HUDSPETH et al., "Sensitivity, Polarity, and Conductance Change in the Response of Vertebrate Hair Cells to Controlled Mechanical Stimuli," <i>Proc. Natl. Acad. Sci. USA</i> , Vol. 74 (6), pp. 2407-2411 (1977)		
APW	AH	JAIN et al., "A Human Recessive Neurosensory Nonsyndromic Hearing Impairment Locus is a Potential Homologue of the Murine Deafness (<i>dn</i>) Locus," <i>Hum. Mol. Genet.</i> , Vol. 4 (12), pp. 2391-2394 (1995)		
APW	AI	KURIMA et al., "Genetic Map Localization of DFNA34 and DFNA36, Two Autosomal Dominant Non-Syndromic Deafness Loci," <i>Am. J. Hum. Genet.</i> , Vol. 67, p. 300, Poster no. 1654 (2000)		
	AJ	KURIMA et al., "Genetic Map Localization of DFNA34 and DFNA36, Two Novel Autosomal Dominant Nonsyndromic Deafness Loci," <i>ARO Abstracts</i>, Vol. 24, p. 266, Abstract no. 949 (2001)		
APW	AK	KURIMA, et al., "Dominant and Recessive Deafness Caused by Mutations of a Novel Gene, <i>TMC1</i> , Required for Cochlear Hair-Cell Function," <i>Nat. Genet.</i> , Vol. 30, pp. 277-284 (2002)		
APW	AL	SCOTT et al., "Refining the DFNB7-DFNB11 Deafness Locus Using Intragenic Polymorphisms in a Novel Gene, <i>TMEM2</i> ," <i>Gene</i> , Vol. 246, pp. 265-274 (2000)		

Examiner Signature		Date Considered	12/7/05
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- * A concise statement of relevance is being submitted in lieu of a translation. 37 CFR 1.98(a)(3).
 + An English-language equivalent/patent, or an English-language abstract, or an English-language version of the search report or action by a foreign patent office in a counterpart foreign application indicating the degree of relevance found by the foreign office is being submitted in lieu of a concise explanation of relevance under 37 CFR 1.98(a)(3).